

# PACAP (6-38), human, ovine, rat

Cat. No.: RP10338

## Overview

|                            |  |
|----------------------------|--|
| <b>Description</b>         | PACAP (6-38) is a potent antagonist of PACAP 38. PACAP (6-38) is much more potent and selective than PACAP (6-27) in the inhibition of PACAP-27-stimulated pituitary adenylate cyclase. $K_i$ values for the inhibition of the enzyme were 7nM and 150 nM, respectively. PACAP (6-38) caused a small but significant (approximately 20%) inhibition of the NANC relaxation due to electrical field stimulation (1 Hz or 10 Hz for 20 s). At these frequencies PACAP (6-38) caused no inhibition of the NANC relaxation in the presence of the P2 purinoceptor antagonist pyridoxal-phosphate-6-azophenyl-2',4'-disulphonic acid (or PPADS) plus the NO-synthase blocker NG-nitro-L-arginine; in preparations pretreated with L-NOARG alone, PACAP (6-38) retained its inhibitory effect. |
| <b>Sequence</b>            | {PHE}{THR}{ASP}{SER}{TYR}{SER}{ARG}{TYR}{ARG}{LYS}{GLN}{MET}{ALA}{VAL}{LYS}{LYS}{TYR}{LEU}{ALA}{ALA}{VAL}{LEU}{GLY}{LYS}{ARG}{TYR}{LYS}{GLN}{ARG}{VAL}{LYS}{ASN}{LYS}-NH <sub>2</sub>  |
| <b>Sequence Shortening</b> | FTDSYSRYRKQMAVKKYLA AVL GKRYKQ RVK NK-NH <sub>2</sub>  |
| <b>Molecular Formula</b>   | C <sub>182</sub> H <sub>300</sub> N <sub>56</sub> O <sub>45</sub> S <sub>1</sub>   |
| <b>C Terminal</b>          | NH <sub>2</sub>  |
| <b>Molecular Weight</b>    | 4024.76  |

## Properties

|                   |  |
|-------------------|--|
| <b>Purity</b>     | > 95%  |
| <b>Solubility</b> | Soluble in water. The contents of this vial have been accurately determined. Both the stopper and the vial have been siliconized. Do not attempt to weigh out a smaller portion of the contents. |
| <b>Form</b>       | Lyophilized  |
| <b>Storage</b>    | Store the peptide at -20°C.  |
| <b>Note</b>       | This peptide is a potent agonist.  |